We claim:

- A method for increasing the yield in glyphosate-resistant
 legumes, which comprises treating the plants with a mixture comprising
 - a) a compound of the formula I

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$$X_m + A$$

I

in which

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x is halogen, C₁-C₄-alkyl or trifluoromethyl,

m is 0 or 1,

20 Q is $C(=CH-CH_3)-COOCH_3$, $C(=CH-OCH_3)-COOCH_3$, $C(=N-OCH_3)-COOCH_3$ or $N(-OCH_3)-COOCH_3$,

A is -O-B, $-CH_2O-B$, $-OCH_2-B$, -CH=CH-B, -C=C-B, $-CH_2O-N=C(R^1)-B$ or $-CH_2O-N=C(R^2)=N-OR^3$, where

B is phenyl, naphthyl, 5-membered or 6-membered hetaryl or 5-membered or 6-membered heterocyclyl, comprising one to three N atoms and/or one O or S atom or one or two O and/or S atoms, the ring systems being unsubstituted or substituted by one to three radicals Ra:

Ra being cyano, nitro, amino, aminocarbonyl, aminothiocarbonyl, halogen, C₁-C₆-alkyl, C₁-C₆-haloalkyl, C₁-C₆-alkylcarbonyl, C₁-C₆-alkylsulfonyl, C₁-C₆-alkylsulfoxyl, C₃-C₆-cycloalkyl, C₁-C₆-alkoxy, C₁-C₆-haloalkoxy,

C₁-C₆-alkyloxycarbonyl, C₁-C₆-alkylthio, C₁-C₆-alkylamino, di-C₁-C₆-alkylamino,

C₁-C₆-alkylaminocarbonyl, di-C₁-C₆-alkylaminocarbonyl, C₁-C₆-alkylaminothiocarbonyl,

 $di-C_1-C_6-alkylaminothiocarbonyl, C_2-C_6-alkenyl,$

45 C₂-C₆-alkenyloxy, phenyl, phenoxy, benzyl,

benzyloxy, 5- or 6-membered heterocyclyl, 5- or 6-membered hetaryl, 5- or 6-membered hetaryloxy, C(=NOR')-OR" or OC(R')2-C(R")=NOR", the cyclic radicals, in turn, being unsubstituted or substituted by one to three radicals Rb:

being cyano, nitro, halogen, amino, Rb aminocarbonyl, aminothiocarbonyl, $C_1-C_6-alkyl$, $C_1-C_6-haloalkyl$, C₁-C₆-alkylsulfonyl, C₁-C₆-alkylsulfoxyl, C_3-C_6 -cycloalkyl, C_1-C_6 -alkoxy, C_1-C_6 -haloalkoxy, C_1-C_6 -alkoxycarbonyl, C_1-C_6 -alkylthio, C_1-C_6 -alkylamino, di-C1-C6-alkylamino, C₁-C₆-alkylaminocarbonyl, di-C₁-C₆-alkylaminocarbonyl, C₁-C₆-alkylaminothiocarbonyl, di-C₁-C₆-alkylaminothiocarbonyl, C_2-C_6 -alkenyl, C_2-C_6 -alkenyloxy, C3-C6-cycloalkyl, C3-C6-cycloalkenyl, phenyl, phenoxy, phenylthio, benzyl, benzyloxy, 5or 6-membered heterocyclyl, 5- or 6-membered hetaryl, 5- or 6-membered hetaryloxy or C(=NOR')-OR''

- R' is hydrogen, cyano, C_1-C_6 -alkyl, C_3-C_6 -cycloalkyl or C_1-C_4 -haloalkyl,
- R" is hydrogen, C₁-C₆-alkyl, C₃-C₆-alkenyl,
 C₃-C₆-alkynyl, C₁-C₄-haloalkyl,
 C₃-C₆-haloalkenyl or C₃-C₆-haloalkynyl,
- R¹ is hydrogen, cyano, C_1-C_4 -alkyl, C_1-C_4 -haloalkyl, C_3-C_6 -cycloalkyl or C_1-C_4 -alkoxy,
- is phenyl, phenylcarbonyl, phenylsulfonyl, 5- or 6-membered hetaryl, 5- or 6-membered hetarylcarbonyl or 5- or 6-membered hetarylsulfonyl, the ring systems being unsubstituted or substituted by one to three radicals Ra,

 C_1 - C_{10} -alkyl, C_3 - C_6 -cycloalkyl, C_2 - C_{10} -alkenyl, C_2 - C_{10} -alkynyl, C_1 - C_{10} -alkylcarbonyl, C_2 - C_{10} -alkynylcarbonyl, C_3 - C_{10} -alkynylcarbonyl, C_1 - C_{10} -alkylsulfonyl or C(R')=NOR", the hydrocarbon radicals of these groups

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being unsubstituted or substituted by one to three radicals R^c:

- being cyano, nitro, amino, aminocarbonyl, RC aminothiocarbonyl, halogen, C1-C6-alkyl, 5 C₁-C₆-haloalkyl, C₁-C₆-alkylsulfonyl, $C_1-C_6-alkylsulfoxyl, C_1-C_6-alkoxy,$ C₁-C₆-haloalkoxy, C₁-C₆-alkoxycarbonyl, C_1-C_6 -alkylthio, C_1-C_6 -alkylamino, di-C₁-C₆-alkylamino, C₁-C₆-alkylaminocarbonyl, 10 di-C₁-C₆-alkylaminocarbonyl, C1-C6-alkylaminothiocarbonyl, di-C1-C6-alkylaminothiocarbonyl, C2-C6-alkenyl, C_2-C_6 -alkenyloxy, C_3-C_6 -cycloalkyl, C₃-C₆-cycloalkyloxy, 5- or 6-membered 15 heterocyclyl, 5- or 6-membered heterocyclyloxy, benzyl, benzyloxy, phenyl, phenoxy, phenylthio, 5- or 6-membered hetaryl, 5- or 6-membered hetaryloxy and hetarylthio, it being possible for the cyclic groups, in turn, to be partially 20 or fully halogenated or to have attached to them one to three radicals Ra, and
 - R^3 is hydrogen, C_1 - C_6 -alkyl, C_2 - C_6 -alkenyl or C_2 - C_6 -alkynyl, the hydrocarbon radicals of these groups being unsubstituted or substituted by one to three radicals R^c ,

and

- 30 b) a glyphosate derivative II in a synergistically active amount.
 - 2. A method as claimed in claim 1, wherein an active ingredient of the formula Ia

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$$O \longrightarrow N \longrightarrow (R^{a'})_{y_i}$$

$$O \longrightarrow N \longrightarrow (R^{b})_{x}$$

$$O \longrightarrow N \longrightarrow (R^{b})_{x}$$

$$O \longrightarrow (R^{b})_{x}$$

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in which T is CH or N and R^a and R^b are halogen or C_1 - C_4 -alkyl, the phenyl group is in the 1- or 5-position and x is 0, 1 or 2 and y is 0 or 1

is used as component a).

- 3. A method as claimed in claim 1 or 2, wherein a fungicidal azole selected from the group consisting of: fluquinconazole, metconazole, prochloraz, propiconazole, prothioconazole, tebuconazole, epoxiconazole or myclobutanil is employed as component a) in addition to the active ingredient of the formula I or Ia.
- 4. A mixture comprising
- a) a compound of the formula I

$$X_m$$

I

٠'n

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in which

- x is halogen, C₁-C₄-alkyl or trifluoromethyl,
- 20 m is 0 or 1,
 - Q is $C(=CH-CH_3)-COOCH_3$, $C(=CH-OCH_3)-COOCH_3$, $C(=N-OCH_3)-COOCH_3$, $C(=N-OCH_3)-COOCH_3$ or $N(-OCH_3)-COOCH_3$,

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A is -O-B, $-CH_2O-B$, $-OCH_2-B$, -CH=CH-B, -C=C-B, $-CH_2O-N=C(R^1)-B$ or $-CH_2O-N=C(R^1)-C(R^2)=N-OR^3$, where

is phenyl, naphthyl, 5-membered or 6-membered hetaryl or 5-membered or 6-membered heterocyclyl, comprising one to three N atoms and/or one O or S atom or one or two O and/or S atoms, the ring systems being unsubstituted or substituted by one to three radicals Ra:

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Ra being cyano, nitro, amino, aminocarbonyl, aminothiocarbonyl, halogen, C₁-C₆-alkyl, C₁-C₆-haloalkyl, C₁-C₆-alkylcarbonyl, C₁-C₆-alkylsulfonyl, C₁-C₆-alkylsulfoxyl, C₃-C₆-cycloalkyl, C₁-C₆-alkoxy, C₁-C₆-haloalkoxy, C₁-C₆-alkyloxycarbonyl, C₁-C₆-alkylthio, C₁-C₆-alkylamino, di-C₁-C₆-alkylamino, di-C₁-C₆-alkylaminocarbonyl, di-C₁-C₆-alkylaminocarbonyl,

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 $C_1-C_6-alkylaminothiocarbonyl,$ $di-C_1-C_6-alkylaminothiocarbonyl,$ $C_2-C_6-alkenyloxy,$ phenyl, phenoxy, benzyl,

benzyloxy, 5- or 6-membered heterocyclyl, 5- or 6-membered hetaryl, 5- or 6-membered hetaryloxy, C(=NOR')-OR'' or $OC(R')_2-C(R'')=NOR''$, the cyclic radicals, in turn, being unsubstituted or substituted by one to three radicals R^b :

RÞ being cyano, nitro, halogen, amino, aminocarbonyl, aminothiocarbonyl, $C_1-C_6-alkyl$, $C_1-C_6-haloalkyl$, $C_1-C_6-alkylsulfonyl$, $C_1-C_6-alkylsulfoxyl$, C_3-C_6 -cycloalkyl, C_1-C_6 -alkoxy, C_1-C_6 -haloalkoxy, C_1-C_6 -alkoxycarbonyl, C_1-C_6 -alkylthio, C_1-C_6 -alkylamino, $di-C_1-C_6-alkylamino$, C₁-C₆-alkylaminocarbonyl, di-C1-C6-alkylaminocarbonyl, C₁-C₆-alkylaminothiocarbonyl, di-C₁-C₆-alkylaminothiocarbonyl, C_2-C_6 -alkenyl, C_2-C_6 -alkenyloxy, C3-C6-cycloalkyl, C3-C6-cycloalkenyl, phenyl, phenoxy, phenylthio, benzyl, benzyloxy, 5or 6-membered heterocyclyl, 5- or 6-membered hetaryl, 5- or 6-membered hetaryloxy or C(=NOR')-OR''

- R' is hydrogen, cyano, C_1-C_6 -alkyl, C_3-C_6 -cycloalkyl or C_1-C_4 -haloalkyl,
- R" is hydrogen, C_1-C_6 -alkyl, C_3-C_6 -alkenyl, C_3-C_6 -alkynyl, C_1-C_4 -haloalkyl, C_3-C_6 -haloalkenyl or C_3-C_6 -haloalkynyl,
- R¹ is hydrogen, cyano, C_1-C_4 -alkyl, C_1-C_4 -haloalkyl, C_3-C_6 -cycloalkyl or C_1-C_4 -alkoxy,
- R² is phenyl, phenylcarbonyl, phenylsulfonyl, 5- or 6-membered hetaryl, 5- or 6-membered hetarylcarbonyl or 5- or 6-membered hetarylsulfonyl, the ring systems being unsubstituted or substituted by one to three radicals R^a,

 C_1-C_{10} -alkyl, C_3-C_6 -cycloalkyl, C_2-C_{10} -alkenyl, C_2-C_{10} -alkynyl, C_1-C_{10} -alkylcarbonyl, C_2-C_{10} -alkenylcarbonyl, C_3-C_{10} -alkynylcarbonyl, C_1-C_{10} -alkylsulfonyl or C(R')=NOR", the hydrocarbon radicals of these groups

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being unsubstituted or substituted by one to three radicals R^c:

- being cyano, nitro, amino, aminocarbonyl, RC aminothiocarbonyl, halogen, C1-C6-alkyl, 5 C₁-C₆-haloalkyl, C₁-C₆-alkylsulfonyl, $C_1-C_6-alkylsulfoxyl, C_1-C_6-alkoxy,$ C_1-C_6 -haloalkoxy, C_1-C_6 -alkoxycarbonyl, C₁-C₆-alkylthio, C₁-C₆-alkylamino, di-C₁-C₆-alkylamino, C₁-C₆-alkylaminocarbonyl, 10 di-C1-C6-alkylaminocarbonyl, C₁-C₆-alkylaminothiocarbonyl, di-C1-C6-alkylaminothiocarbonyl, C2-C6-alkenyl, C2-C6-alkenyloxy, C3-C6-cycloalkyl, C₃-C₆-cycloalkyloxy, 5- or 6-membered 15 heterocyclyl, 5- or 6-membered heterocyclyloxy, benzyl, benzyloxy, phenyl, phenoxy, phenylthio, 5- or 6-membered hetaryl, 5- or 6-membered hetaryloxy and hetarylthio, it being possible for the cyclic groups, in turn, to be partially 20 or fully halogenated or to have attached to them one to three radicals Ra, and
 - R^3 is hydrogen, C_1 - C_6 -alkyl or C_2 - C_6 -alkenyl or C_2 - C_6 -alkynyl, the hydrocarbon radicals of these groups being unsubstituted or substituted by one to three radicals R^c ,

and

30 b) a glyphosate derivative II

in such an amount that the yield is increased synergistically.

- 35 5. A mixture as claimed in claim 4, comprising
 - a) pyraclostrobin and
 - b) a glyphosate derivative II.
- 40 6. A mixture as claimed in claim 5, wherein component a) comprises an azole selected from the group consisting of: metconazole, myclobutanil, epoxiconazole, propiconazole, prothioconazole and tebuconazole in addition to the active ingredient pyraclostrobin.

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